

# Richard K Haynes

## List of Publications by Year in descending order

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193  
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7,322  
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57758

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74163

75  
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219  
all docs

219  
docs citations

219  
times ranked

4875  
citing authors

#	ARTICLE	IF	CITATIONS
1	Artemisinins: their growing importance in medicine. Trends in Pharmacological Sciences, 2008, 29, 520-527.	8.7	301
2	A single amino acid residue can determine the sensitivity of SERCAs to artemisinins. Nature Structural and Molecular Biology, 2005, 12, 628-629.	8.2	232
3	Artemisone—A Highly Active Antimalarial Drug of the Artemisinin Class. Angewandte Chemie - International Edition, 2006, 45, 2082-2088.	13.8	222
4	From Qinghao, Marvelous Herb of Antiquity, to the Antimalarial Trioxane Qinghaosu and Some Remarkable New Chemistry. Accounts of Chemical Research, 1997, 30, 73-79.	15.6	209
5	From Artemisinin to New Artemisinin Antimalarials: Biosynthesis, Extraction, Old and New Derivatives, Stereochemistry and Medicinal Chemistry Requirements. Current Topics in Medicinal Chemistry, 2006, 6, 509-537.	2.1	208
6	Possible modes of action of the artemisinin-type compounds. Trends in Parasitology, 2001, 17, 122-126.	3.3	207
7	Artemisinins. Postgraduate Medical Journal, 2005, 81, 71-78.	1.8	200
8	Artemisinins: mechanisms of action and potential for resistance. Drug Resistance Updates, 2004, 7, 233-244.	14.4	180
9	Highly Antimalaria-Active Artemisinin Derivatives: Biological Activity Does Not Correlate with Chemical Reactivity. Angewandte Chemie - International Edition, 2004, 43, 1381-1385.	13.8	137
10	Artemisinin and derivatives: the future for malaria treatment?. Current Opinion in Infectious Diseases, 2001, 14, 719-726.	3.1	117
11	Artemisinins Inhibit Trypanosoma cruzi and Trypanosoma brucei rhodesiense In Vitro Growth. Antimicrobial Agents and Chemotherapy, 2007, 51, 1852-1854.	3.2	116
12	Neurotoxic Mode of Action of Artemisinin. Antimicrobial Agents and Chemotherapy, 2002, 46, 821-827.	3.2	111
13	The Fe <sup>2+</sup> -Mediated Decomposition, PfATP6 Binding, and Antimalarial Activities of Artemisone and Other Artemisinins: The Unlikelihood of Centered Radicals as Bioactive Intermediates. ChemMedChem, 2007, 2, 1480-1497.	3.2	107
14	Highly Enantioselective Phenyl Transfer to Aryl Aldehydes Catalyzed by Easily Accessible Chiral Tertiary Aminonaphthol. Journal of Organic Chemistry, 2005, 70, 1093-1095.	3.2	106
15	New reactions of triplet oxygen which avoid the spin barrier. Journal of the Chemical Society Perkin Transactions 1, 1975, , 2055.	0.9	98
16	Artemisinins: activities and actions. Microbes and Infection, 2004, 6, 1339-1346.	1.9	95
17	First Assessment in Humans of the Safety, Tolerability, Pharmacokinetics, and Ex Vivo Pharmacodynamic Antimalarial Activity of the New Artemisinin Derivative Artemisone. Antimicrobial Agents and Chemotherapy, 2008, 52, 3085-3091.	3.2	90
18	Artesunate and Dihydroartemisinin (DHA): Unusual Decomposition Products Formed under Mild Conditions and Comments on the Fitness of DHA as an Antimalarial Drug. ChemMedChem, 2007, 2, 1448-1463.	3.2	86

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19	In vitro anti-cancer effects of artemisone nano-vesicular formulations on melanoma cells. <i>Nanomedicine: Nanotechnology, Biology, and Medicine</i> , 2015, 11, 2041-2050.	3.3	86
20	The behaviour of qinghaosu (artemisinin) in the presence of heme iron(II) and (III). <i>Tetrahedron Letters</i> , 1996, 37, 253-256.	1.4	85
21	Antimalarial efficacy and drug interactions of the novel semi-synthetic endoperoxide artemisone in vitro and in vivo. <i>Journal of Antimicrobial Chemotherapy</i> , 2007, 59, 658-665.	3.0	83
22	Re-evaluation of how artemisinins work in light of emerging evidence of in vitro resistance. <i>Trends in Molecular Medicine</i> , 2006, 12, 200-205.	6.7	82
23	The behaviour of qinghaosu (artemisinin) in the presence of non-heme iron(II) and (III). <i>Tetrahedron Letters</i> , 1996, 37, 257-260.	1.4	76
24	Facile Oxidation of Leucomethylene Blue and Dihydroflavins by Artemisinins: Relationship with Flavoenzyme Function and Antimalarial Mechanism of Action. <i>ChemMedChem</i> , 2010, 5, 1282-1299.	3.2	76
25	Reaction of Metallated tert-Butyl(phenyl)phosphane Oxide with Electrophiles as a Route to Functionalized Tertiary Phosphane Oxides: Alkylation Reactions. <i>European Journal of Organic Chemistry</i> , 2000, 2000, 3205-3216.	2.4	75
26	C-10 Ester and Ether Derivatives of Dihydroartemisinin $\hat{\sim}$ 10- $\hat{I}$ $\pm$ Artesunate, Preparation of Authentic 10- $\hat{I}$ $^2$ Artesunate, and of Other Ester and Ether Derivatives Bearing Potential Aromatic Intercalating Groups at C-10. <i>European Journal of Organic Chemistry</i> , 2002, 2002, 113-132.	2.4	74
27	Artemisone and Artemiside Control Acute and Reactivated Toxoplasmosis in a Murine Model. <i>Antimicrobial Agents and Chemotherapy</i> , 2009, 53, 4450-4456.	3.2	74
28	Considerations on the Mechanism of Action of Artemisinin Antimalarials: Part 1 - The $\hat{\&\#39}$ Carbon Radical $\hat{\&\#39}$ ; and $\hat{\&\#39}$ Heme $\hat{\&\#39}$ ; Hypotheses. <i>Infectious Disorders - Drug Targets</i> , 2014, 13, 217-277.	0.8	72
29	Ring opening of artemisinin (qinghaosu) and dihydroartemisinin and interception of the open hydroperoxides with Formation of N-oxides $\hat{\&\#39}$ a chemical model for antimalarial mode of action. <i>Tetrahedron Letters</i> , 1999, 40, 4715-4718.	1.4	71
30	Differential effects on angiogenesis of two antimalarial compounds, dihydroartemisinin and artemisone: Implications for embryotoxicity. <i>Toxicology</i> , 2007, 241, 66-74.	4.2	68
31	Artemisinin Antimalarials Do Not Inhibit Hemozoin Formation. <i>Antimicrobial Agents and Chemotherapy</i> , 2003, 47, 1175-1175.	3.2	67
32	Anticancer Properties of Distinct Antimalarial Drug Classes. <i>PLoS ONE</i> , 2013, 8, e82962.	2.5	67
33	A novel endoperoxide and related sesquiterpenes from <i>Artemisia annua</i> which are possibly derived from allylic hydroperoxides. <i>Tetrahedron</i> , 1998, 54, 4345-4356.	1.9	65
34	Interactions between Artemisinins and other Antimalarial Drugs in Relation to the Cofactor Model $\hat{\&\#39}$ A Unifying Proposal for Drug Action. <i>ChemMedChem</i> , 2012, 7, 2204-2226.	3.2	63
35	Recent progress in the development of anti-malarial quinolones. <i>Malaria Journal</i> , 2014, 13, 339.	2.3	63
36	Preparation of stable, camphor-derived, optically active allyl and alkyl sulfoxides and thermal epimerization of the allyl sulfoxides. <i>Journal of Organic Chemistry</i> , 1988, 53, 2881-2889.	3.2	62

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37	Artemisone effective against murine cerebral malaria. <i>Malaria Journal</i> , 2010, 9, 227.	2.3	62
38	Preparation of Enantiomerically Pure Tertiary Phosphine Oxides from, and Assay of Enantiomeric Purity with, (Rp)- and (Sp)-tert-Butylphenylphosphinothioic Acids. <i>Journal of Organic Chemistry</i> , 1994, 59, 2919-2921.	3.2	55
39	Catalysed oxygenation of allylic hydroperoxides derived from qinghao (artemisinic) acid. Conversion of qinghao acid into dehydroqinghaosu (artemisitenone) and qinghaosu (artemisinin). <i>Journal of the Chemical Society Chemical Communications</i> , 1990, , 451.	2.0	54
40	Use of .beta.-sulfonyl vinyl ketones as equivalents to vinyl ketones in the Robinson annelation. Convergent, highly stereoselective preparation of a hydrindanol related to vitamin D from 2-methylcyclopent-2-enone and lithiated (E)-but-2-enyldiphenylphosphine oxide. <i>Journal of Organic Chemistry</i> , 1989, 54, 5162-5170.	3.2	52
41	Aprotic conjugate addition of allyllithium reagents bearing polar groups to cyclic enones. 1. 3-Alkylallyl systems. <i>Journal of the American Chemical Society</i> , 1988, 110, 5411-5423.	13.7	51
42	Stereoselective preparation of functionalized tertiary P-chiral phosphine oxides by nucleophilic addition of lithiated tert-butylphenylphosphine oxide to carbonyl compounds. <i>Tetrahedron Letters</i> , 1996, 37, 4729-4732.	1.4	49
43	Hexamethylphosphoramide-mediated conjugate addition of (alkylthio)-, (phenylthio)-, and (phenylseleno)allyllithium reagents to 2-cyclopentenone. <i>Journal of Organic Chemistry</i> , 1981, 46, 3790-3795.	3.2	48
44	Reactions of Antimalarial Peroxides with Each of Leucomethylene Blue and Dihydroflavins: Flavin Reductase and the Cofactor Model Exemplified. <i>ChemMedChem</i> , 2011, 6, 279-291.	3.2	47
45	Reactions of (RP)- and (SP)-tert-butylphenylphosphinobromidates and tert-butylphenylthionophosphinochloridates with heteroatom nucleophiles; preparation of P-chiral binol phosphinates and related compounds. <i>Tetrahedron Letters</i> , 2001, 42, 453-456.	1.4	46
46	Air-stable P-stereogenic secondary phosphine oxides as chiral monodentate ligands for asymmetric catalytic carbon-carbon bond formation. <i>Tetrahedron: Asymmetry</i> , 2003, 14, 2821-2826.	1.8	46
47	In vitro study of the anti-cancer effects of artemisone alone or in combination with other chemotherapeutic agents. <i>Cancer Chemotherapy and Pharmacology</i> , 2011, 67, 569-577.	2.3	46
48	Copper(II) Trifluoromethanesulfonate-Induced Cleavage Oxygenation of Allylic Hydroperoxides Derived from Qinghao Acid in the Synthesis of Qinghaosu Derivatives: Evidence for the Intermediacy of Enols. <i>Journal of the American Chemical Society</i> , 1995, 117, 11098-11105.	13.7	45
49	Dihydroartemisinin inhibits the human erythroid cell differentiation by altering the cell cycle. <i>Toxicology</i> , 2012, 300, 57-66.	4.2	45
50	Low-Temperature X-ray Crystal-Structure Analysis of the Thermally Unstable Lithiated 2-Butenyltert-Butyl Sulfide: A comparison with modelab initio MO calculations. <i>Helvetica Chimica Acta</i> , 1988, 71, 299-311.	1.6	44
51	Glucocorticosteroids in Nano-Sterically Stabilized Liposomes Are Efficacious for Elimination of the Acute Symptoms of Experimental Cerebral Malaria. <i>PLoS ONE</i> , 2013, 8, e72722.	2.5	41
52	Formation of iodides and esters from alcohols and tributyl-diiodophosphorane and diiodotriphenylphosphorane. <i>Australian Journal of Chemistry</i> , 1982, 35, 517.	0.9	40
53	Absorption of the novel artemisinin derivatives artemisone and artemiside: Potential application of Pheroida,® technology. <i>International Journal of Pharmaceutics</i> , 2011, 414, 260-266.	5.2	40
54	Novel formation of isomeric bicyclo[3.2.0]heptan-1-ols from phenyl vinyl sulfoxide and the cyclopentanone lithium enolate generated by conjugate addition of lithiated (E)-but-2-enyldiphenylphosphine oxide to 2-methylcyclopent-2-enone. <i>Journal of Organic Chemistry</i> , 1991, 56, 5785-5790.	3.2	39

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55	Simultaneous determination of artemether and its major metabolite dihydroartemisinin in plasma by gas chromatography–mass spectrometry-selected ion monitoring. <i>Biomedical Applications</i> , 1999, 731, 251-260.	1.7	39
56	Stereoselective Preparation of 10 <sup>1±</sup> - and 10 <sup>12</sup> -Aryl Derivatives of Dihydroartemisinin. <i>European Journal of Organic Chemistry</i> , 2003, 2098-2114.	2.4	39
57	The Artemisinin Derivative Artemisone Is a Potent Inhibitor of Human Cytomegalovirus Replication. <i>Antimicrobial Agents and Chemotherapy</i> , 2018, 62, .	3.2	39
58	Artemisone and Artemiside Are Potent Panreactive Antimalarial Agents That Also Synergize Redox Imbalance in Plasmodium falciparum Transmissible Gametocyte Stages. <i>Antimicrobial Agents and Chemotherapy</i> , 2018, 62, .	3.2	39
59	HMPA-Mediated conjugate addition of alkyl- and phenylthioallyl anions to cyclopentenone. <i>Tetrahedron Letters</i> , 1980, 21, 573-576.	1.4	38
60	Extraction of artemisinin and artemisinic acid: preparation of artemether and new analogues. <i>Transactions of the Royal Society of Tropical Medicine and Hygiene</i> , 1994, 88, 23-26.	1.8	38
61	Comparative <i>Ex Vivo</i> Activity of Novel Endoperoxides in Multidrug-Resistant Plasmodium falciparum and P. vivax. <i>Antimicrobial Agents and Chemotherapy</i> , 2012, 56, 5258-5263.	3.2	38
62	Direct formation of 3,3,6,6-Tetraaryl-1,2-dioxans from 1,1-Diarylethylenes and oxygen, catalysed by antimony(V) chloride. <i>Australian Journal of Chemistry</i> , 1978, 31, 1737.	0.9	36
63	Radical mechanism of action of the artemisinin-type compounds. <i>Trends in Parasitology</i> , 2001, 17, 266-267.	3.3	36
64	Chiral Bisphosphinite Metalloligands Derived from a P-Chiral Secondary Phosphine Oxide. <i>Inorganic Chemistry</i> , 2004, 43, 4921-4926.	4.0	36
65	Convenient Access Both to Highly Antimalaria-Active 10-Arylaminoartemisinins, and to 10-Alkyl Ethers Including Artemether, Arteether, and Artelinate. <i>ChemBioChem</i> , 2005, 6, 659-667.	2.6	36
66	Aprotic conjugate addition of allyllithium reagents bearing polar groups to cyclic enones. 2. 2-Alkyl-, 2,3-dialkyl- and 1,3-dialkylallyl systems. <i>Journal of the American Chemical Society</i> , 1988, 110, 5423-5433.	13.7	34
67	Preparation of <i>N</i> -Sulfonyl- and <i>N</i> -Carbonyl-1 <sup>±</sup> -Azaartemisinins with Greatly Enhanced Thermal Stabilities: <i>in vitro</i> Antimalarial Activities. <i>ChemMedChem</i> , 2007, 2, 1464-1479.	3.2	34
68	Efficient Preparation of Novel Qinghaosu (Artemisinin) Derivatives: Conversion of Qinghao (Artemisinic) Acid into Deoxoqinghaosu Derivatives and 5-Carba-4-deoxoartesunic Acid. <i>Synlett</i> , 1992, 1992, 481-483.	1.8	32
69	Some further observations on the Lewis-acid-catalysed oxygenation of ergosteryl acetate. <i>Australian Journal of Chemistry</i> , 1978, 31, 121.	0.9	31
70	An Improved Method for the Isolation of Qinghao (Artemisinic) Acid from <i>Artemisia annua</i> . <i>Planta Medica</i> , 1993, 59, 562-563.	1.3	31
71	<i>Neospora caninum</i> : In vivo and in vitro treatment with artemisone. <i>Veterinary Parasitology</i> , 2012, 187, 99-104.	1.8	31
72	Iron(III) and copper(II) catalysed transformations of fatty acid hydroperoxides: efficient generation of peroxy radicals with copper(II) trifluoromethanesulphonate. <i>Journal of the Chemical Society Chemical Communications</i> , 1990, , 1102.	2.0	30

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73	Highly Diastereoselective Conjugate Addition of Lithiated $\hat{1}^3$ -Crotonolactone (But-2-en-4-olide) to Cyclic Enones To Give Syn-Adducts: Application to a Brefeldin Synthesis. <i>Journal of Organic Chemistry</i> , 1997, 62, 4552-4553.	3.2	30
74	Accessible and distinct decoquinane derivatives active against <i>Mycobacterium tuberculosis</i> and apicomplexan parasites. <i>Communications Chemistry</i> , 2018, 1, .	4.5	30
75	Lewis-acid-catalysed oxygenation of 1,1'-bicyclohexenyl and $\hat{1}^3$ -terpinene. <i>Reactions in dichloromethane and liquid sulphur dioxide. Australian Journal of Chemistry</i> , 1978, 31, 131.	0.9	29
76	Preparation of a Bicyclic Analog of Qinghao (Artemisinic) Acid via a Lewis Acid Catalyzed Ionic Diels-Alder Reaction Involving a Hydroxy Diene and Cyclic Enone and Facile Conversion into (.+)-6,9-Desdimethylqinghaosu. <i>Journal of Organic Chemistry</i> , 1994, 59, 4743-4748.	3.2	29
77	Interaction of Artemisinins with Oxyhemoglobin Hb $\hat{Fe}^{II}$ , Hb $\hat{Fe}^{II}$ , CarboxyHb $\hat{Fe}^{II}$ , Heme $\hat{Fe}^{II}$ , and Carboxyheme $\hat{Fe}^{II}$ : Significance for Mode of Action and Implications for Therapy of Cerebral Malaria. <i>ChemMedChem</i> , 2009, 4, 2045-2053.	3.2	29
78	Reply to Comments on 'Highly Antimalaria-Active Artemisinin Derivatives: Biological Activity Does Not Correlate with Chemical Reactivity?'. <i>Angewandte Chemie - International Edition</i> , 2005, 44, 2064-2065.	13.8	28
79	Synthesis of Artemiside and Its Effects in Combination with Conventional Drugs against Severe Murine Malaria. <i>Antimicrobial Agents and Chemotherapy</i> , 2012, 56, 163-173.	3.2	28
80	Synthesis, in vitro antimalarial activities and cytotoxicities of amino-artemisinin-ferrocene derivatives. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2018, 28, 289-292.	2.2	28
81	A new resolution procedure for the preparation of both (R)-(+)- and (S)-(-)-4-tert-butoxycyclopent-2-enone from racemic 4-tert-butoxycyclopent-2-enone and conversion of (R)-(+)-4-tert-butoxycyclopent-2-enone into (R)-(+)-4-acetoxycyclopent-2-enone. A new method for the determination of the enantiomeric purities of the resolved enones. <i>Journal of Organic Chemistry</i> , 1991, 56, 4760-4766.	3.2	27
82	Repurposing of antiparasitic drugs: the hydroxy-naphthoquinone buparvaquone inhibits vertical transmission in the pregnant neosporosis mouse model. <i>Veterinary Research</i> , 2016, 47, 32.	3.0	27
83	Kinetically controlled, stereoselective formation of vinylic sulfones by conjugate addition of lithiated 3-allylallylic sulfones to cyclic enones. <i>Journal of Organic Chemistry</i> , 1989, 54, 1960-1968.	3.2	26
84	Preparation of bi- and tridentate doubly P-chiral diphosphine dioxide ligands for asymmetric catalysis. <i>Tetrahedron Letters</i> , 1996, 37, 4733-4736.	1.4	26
85	Completely stereoselective $\hat{P}i-C$ bond formation via base-induced [1,3]- and [1,2]-intramolecular rearrangements of aryl phosphinates, phosphinoamidates and related compounds: generation of P-chiral $\hat{1}^2$ -hydroxy, $\hat{1}^2$ -mercapto- and $\hat{1}^2$ -amino tertiary phosphine oxides and phosphine sulfides. <i>Tetrahedron Letters</i> , 2001, 42, 457-460.	1.4	26
86	A Partial Convergence in Action of Methylene Blue and Artemisinins: Antagonism with Chloroquine, a Reversal with Verapamil, and an Insight into the Antimalarial Activity of Chloroquine. <i>ChemMedChem</i> , 2011, 6, 1603-1615.	3.2	26
87	Synthesis, antimalarial activities and cytotoxicities of amino-artemisinin-1,2-disubstituted ferrocene hybrids. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2018, 28, 3161-3163.	2.2	26
88	Establishment of an In Vitro screening model for neurodegeneration induced by antimalarial drugs of the artemisinin-type. <i>Neurotoxicity Research</i> , 2000, 2, 37-49.	2.7	25
89	Expression in Yeast Links Field Polymorphisms in PfATP6 to In Vitro Artemisinin Resistance and Identifies New Inhibitor Classes. <i>Journal of Infectious Diseases</i> , 2013, 208, 468-478.	4.0	25
90	Evaluation of Artemisone Combinations in <i>Aotus</i> Monkeys Infected with <i>Plasmodium falciparum</i> . <i>Antimicrobial Agents and Chemotherapy</i> , 2009, 53, 3592-3594.	3.2	24

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91	The diastereospecific aprotic conjugate addition reactions of carbanions derived from allylic sulfoxides and allylic phosphine oxides.. Tetrahedron Letters, 1985, 26, 1565-1568.	1.4	23
92	In vitro skin permeation of artemisone and its nano-vesicular formulations. International Journal of Pharmaceutics, 2016, 503, 1-7.	5.2	23
93	Topical Delivery of Artemisone, Clofazimine and Decoquinat Encapsulated in Vesicles and Their In vitro Efficacy Against Mycobacterium tuberculosis. AAPS PharmSciTech, 2019, 20, 33.	3.3	23
94	In vitro effects of new artemisinin derivatives in Neospora caninum-infected human fibroblasts. International Journal of Antimicrobial Agents, 2015, 46, 88-93.	2.5	22
95	In vitro activity of artemisone and artemisinin derivatives against extracellular and intracellular Helicobacter pylori. International Journal of Antimicrobial Agents, 2016, 48, 101-105.	2.5	22
96	Artemisone demonstrates synergistic antiviral activity in combination with approved and experimental drugs active against human cytomegalovirus. Antiviral Research, 2019, 172, 104639.	4.1	22
97	Lewis acid catalysed oxygenation of ergosteryl acetate by triplet oxygen. Journal of the Chemical Society Chemical Communications, 1974, , 511.	2.0	21
98	The First Examples of Enantiomerically Pure Diphosphane Dioxides <sup>2+</sup> (RP,RP)- and (SP,SP)-1,2-Di-ter-butyl-1,2-diphenyldiphosphane 1,2-Dioxides, and (RP)- and (SP)-1-tert-Butyl-1,2,2-triphenyldiphosphane 1,2-Dioxides. Chemistry - A European Journal, 1997, 3, 2052-2057.	3.3	21
99	The evaluation of the anti-cancer drug elesclomol that forms a redox-active copper chelate as a potential anti-tubercular drug. IUBMB Life, 2019, 71, 532-538.	3.4	21
100	Die Oxidation von Orcin mit K <sub>3</sub> [Fe(CN) <sub>6</sub> ] im Strömungsrohr. Chemische Berichte, 1974, 107, 3733-3748.	0.2	20
101	The diastereospecific aprotic conjugate addition reactions of allylic anions-mechanistic aspects.. Tetrahedron Letters, 1985, 26, 1569-1572.	1.4	20
102	Preparation of stable, camphor-derived, optically active allylic sulfoxides. Tetrahedron Letters, 1985, 26, 6381-6384.	1.4	20
103	Methylene Homologues of Artemisone: An Unexpected Structure-Activity Relationship and a Possible Implication for the Design of C10-Substituted Artemisinins. ChemMedChem, 2016, 11, 1469-1479.	3.2	20
104	The preparation of (Methylthio)- and (Methylseleno)-tri(alkyl or aryl)phosphonium salts and their reactions with carboxylic acids and alcohols. Australian Journal of Chemistry, 1984, 37, 1183.	0.9	18
105	The preparation of D-ring-contracted analogues of Qinghaosu (Artemisinin) from Qinghao (Artemisinic) acid and their In vitro activity against Plasmodium falciparum. Tetrahedron Letters, 1995, 36, 4641-4642.	1.4	18
106	Evaluation and optimization of synthetic routes from dihydroartemisinin to the alkylamino-artemisinins artemiside and artemisone: A test of N-glycosylation methodologies on a lipophilic peroxide. Tetrahedron, 2018, 74, 5156-5171.	1.9	18
107	Synthetic utilization of highly stereoselective conjugate addition reactions of phosphorus and sulfur stabilized allylic carbanions. Pure and Applied Chemistry, 1993, 65, 647-654.	1.9	18
108	The preparation of (R)- and (S)-(E)-but-2-enyl-t-butylphenylphosphine oxides and their enantiospecific conversion into enantiomeric hydrindenones related to vitamin D. Journal of the Chemical Society Chemical Communications, 1991, , 58.	2.0	17



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109	Artemisinin and Heme. <i>Antimicrobial Agents and Chemotherapy</i> , 2003, 47, 2712-2713.	3.2	17
110	Treatment of Murine Cerebral Malaria by Artemisone in Combination with Conventional Antimalarial Drugs: Antiplasmodial Effects and Immune Responses. <i>Antimicrobial Agents and Chemotherapy</i> , 2014, 58, 4745-4754.	3.2	17
111	Straightforward conversion of decoquinatate into inexpensive tractable new derivatives with significant antimalarial activities. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2016, 26, 3006-3009.	2.2	17
112	Activities of 11-aza-artemisinin and <i>N</i> -sulfonyl Derivatives against Asexual and Transmissible Malaria Parasites. <i>ChemMedChem</i> , 2017, 12, 2086-2093.	3.2	17
113	An in vitro ADME and in vivo Pharmacokinetic Study of Novel TB-Active Decoquinatate Derivatives. <i>Frontiers in Pharmacology</i> , 2019, 10, 120.	3.5	17
114	Elimination of <i>Schistosoma mansoni</i> in infected mice by slow release of artemisone. <i>International Journal for Parasitology: Drugs and Drug Resistance</i> , 2017, 7, 241-247.	3.4	16
115	Preliminary Evaluation of Artemisinin-Cholesterol Conjugates as Potential Drugs for the Treatment of Intractable Forms of Malaria and Tuberculosis. <i>ChemMedChem</i> , 2018, 13, 67-77.	3.2	16
116	Formulation of Natural Oil Nano-Emulsions for the Topical Delivery of Clofazimine, Artemisone and Decoquinatate. <i>Pharmaceutical Research</i> , 2018, 35, 186.	3.5	16
117	Optimal 10-Aminoartemisinins With Potent Transmission-Blocking Capabilities for New Artemisinin Combination Therapies' Activities Against Blood Stage <i>P. falciparum</i> Including PfK13 C580Y Mutants and Liver Stage <i>P. berghei</i> Parasites. <i>Frontiers in Chemistry</i> , 2019, 7, 901.	3.6	16
118	A Drug Repurposing Approach for Antimalarials Interfering with SARS-CoV-2 Spike Protein Receptor Binding Domain (RBD) and Human Angiotensin-Converting Enzyme 2 (ACE2). <i>Pharmaceuticals</i> , 2021, 14, 954.	3.8	16
119	A route to prostaglandin precursors from 1-(phenylthio)-2-octenyllithium.. <i>Tetrahedron Letters</i> , 1985, 26, 3385-3388.	1.4	15
120	Facile Preparation of N-Glycosylated 10-Piperazinyl Artemisinin Derivatives and Evaluation of Their Antimalarial and Cytotoxic Activities. <i>Molecules</i> , 2018, 23, 1713.	3.8	15
121	The Case for Development of 11-Aza-artemisinins for Malaria. <i>Current Medicinal Chemistry</i> , 2015, 22, 3607-3630.	2.4	15
122	An Extremely Simple Route to a Prostaglandin Precursor: Hexamethylphosphoric-Triamide-Mediated Conjugate Addition of 1-(Phenylthio)Oct-2-Enyllithium to 4-Tert-Butoxycyclopent-2-Enone and Triphenyltin-Chloride-Assisted Reaction of the Enolate With Methyl 7-Iodohept-5-Ynoate. <i>Australian Journal of Chemistry</i> , 1987, 40, 1211.	0.9	14
123	Preparation of hydrindenones from 2-methylcyclopent-2-enone and the carbanion of (E)-but-2-enyldiphenylphosphine oxide: efficient enolate trapping with $\text{I}^2$ -sulphonylvinyl ketones. <i>Journal of the Chemical Society Chemical Communications</i> , 1987, , 92-94.	2.0	14
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